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(FILE 'HOME' ENTERED AT 16:09:18 ON 08 AUG 2004)

FILE 'REGISTRY' ENTERED AT 16:09:28 ON 08 AUG 2004 E LIMN204/MF

L1 1 S E3

FILE 'CAPLUS' ENTERED AT 16:10:15 ON 08 AUG 2004

L2 170 S L1 AND PREPARATION

L3 0 S L2 AND OXYGEN AND INERT

L4 27 S L2 AND (AIR OR OXYGEN)

L5 5 S L4 AND (ARGON OR NITROGEN)

FILE 'REGISTRY' ENTERED AT 16:49:51 ON 08 AUG 2004

E LI203/MF

L6 1 S E3

FILE 'CAPLUS' ENTERED AT 16:51:02 ON 08 AUG 2004

L7 27 S L6

L8 7 S L7 AND (MANGANESE OR MN)

L8 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2003:117764 CAPLUS

DOCUMENT NUMBER:

138:157658

TITLE:

Alumina-based glass-ceramics with high hardness for

ADDLICATION NO

DATE

use as abrasives

INVENTOR(S):

Rosenflanz, Anatoly Z.

PATENT ASSIGNEE(S):

3M Innovative Properties Company, USA

SOURCE:

PCT Int. Appl., 112 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

1Ω

KIND

PATENT INFORMATION:

DATENT NO

| | PAT | ENT I | NO. | | | | | DATE | | | APPL: | ICAT | ION I | | | | ATE | | |
|-------|------|-------|--|---|--|---|--|--|--|--|---|---|---|---|---|---|---|---|----|
| | | | 01178 01178 | 34 | | A2 C1 | | 2003 2003 | 0213 | 1 | NO 21 | 002- | US24 | | | | 00208 | 302 | |
| | | W: | AE. CO. GM. LS. PL. UA. GH. CH. | AG. CR, HR. LT. PT. UG. GM. CY. SE. | AL. CU. HU. LU. RO. UZ. KE. CZ. | AM. CZ. ID. LV. RU. VN. LS. DE. TR. | AT DE IL MA SD YU MW DK | . AU. . DK. . IN. . MD. . SE. . ZA. . MZ. . EE. | AZ. DM. IS. MG. SG. ZM. SD. ES. | BA. DZ. JP. MK. SI. ZW. SL. FI. | EC. KE. MN. SK. AM. SZ. FR. | EE. KG. MW. SL. AZ. TZ. GB. | ES. KP. MX. TJ. BY. UG. GR. | FI. KR. MZ. TM. KG. ZM. IE. | GB, KZ, NO, TN, KZ, ZW, IT, | GD, LC. NZ. TR. MD. AT. LU. | GE, LK, OM, TT, RU, BE, MC, | GH, LR. PH, TZ, TJ, BG, NL, | ТМ |
| | EP : | 1432 | 659 | | | | | 2004 | 0630 | I | EP 20 | 002- | 7504 | 13 | | 20 | 30208 | 302 | |
| | | R: | | | | | | . ES. . RO. | | | | | | | | | MC, | PT, | |
| | EP : | 1432 | | | | | | 2004 | | | | | | | | | 0208 | 302 | |
| | | R: | | | | | | . ES. . RO. | | | | | | | | | MC, | PT, | |
| | RR : | 2002 | | | | | | 2004 | | | | | | | | | 00208 | 302 | |
| PRIOR | | | | | | • | | | | | | | 9225 | | | | 00108 | 302 | |
| | | | | | | | | | | | | | 9225 | | | A 20 | 00108 | 302 | |
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| | 0.3 | | | | _ | | | | | | | | | | V | | | | |
| AB | Glas | SS-C | eram | ICS T | tor (| use i | as a | abras | ive i | part | ıcles | s wi | th h | 1an i | nardr | าess | con | tain | |

AB Glass-ceramics for use as abrasive particles with high hardness contain alumina and other oxides with less than 10 weight% in As203. B203. Ge02. P205, Si02, Te02, and V205 combined and are formed by heat-treatment of the glass compns. into glass-ceramics. The glass-ceramics also contain other components than alumina (such as La203, Zr02, Al, Y203, Gd203, Mg. Mg0, Ti02, CaF2, Nb205. Ta205, Sr0, Mn203, Cr203 and/or Ce02) and may be crushed to form the abrasive particles. The abrasive particles can be incorporated into a variety of abrasive articles such as bonded abrasives.

coated abrasives, nonwoven abrasives and abrasive brushes. IT Brushes (abrasives: aluminate glass ceramic hardness abrasive steel substrate) Abrasion Glass ceramics Glass transition temperature Grain size Hardness (mechanical) Heat treatment (aluminate glass ceramic hardness abrasive steel substrate) IT Size reduction (crushing, of glass-ceramics; aluminate glass ceramic hardness abrasive steel substrate) IT Abrasives (particles; aluminate glass ceramic hardness abrasive steel substrate) IT Galvanized steel RL: NUU (Other use, unclassified): USES (Uses) (substrates: aluminate glass ceramic hardness abrasive steel substrate) 12003-65-5. Aluminum lanthanum oxide (AlLaO3) 12005-21-9, Aluminum 12043-90-2, Aluminum lanthanum oxide yttrium oxide (Al5Y3012) 116590-85-3. Yttrium zirconium oxide (Y0.15Zr0.8501.93) (AlllLa018) 140418-71-9, Cerium zirconium oxide (Ce0.6Zr0.402) RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative) (crystallized phase; aluminate glass ceramic hardness abrasive steel substrate) 144-55-8. Sodium carbonate (NaHCO3), processes 1302-74-5. Corundum, IT 1303-86-2, Boron oxide (B203), processes 1305-78-8, Calcium oxide (CaO), processes 1306-38-3, Ceria, processes 1308-38-9, Chromium oxide (Cr203), processes 1308-87-8. Dysprosium oxide (Dy203) 1308-96-9. Europium oxide (Eu203) 1309-37-1. Ferric oxide. processes 1309-48-4. Magnesium oxide (MgO), processes 1310-53-8, Germanium oxide (GeO2), processes 1312-81-8, Lanthanum oxide (La2O3) 1313-96-8. 1313-97-9, Neodymium oxide (Nd203) 1314-11-0. Niobium oxide (Nb205) Strontium oxide (SrO), processes 1314-23-4, Zirconium oxide (ZrO2), 1314-36-9, Yttrium oxide (Y2O3), processes Phosphorus oxide (P205), processes 1314-61-0, Tantalum oxide (Ta205) 1314-62-1, Vanadium oxide (V205), processes 1317-34-6, Manganese 1327-53-3, Arsenic oxide (As203) 1344-28-1, Aluminum oxide (Mn203) 7429-90-5, Aluminum, processes oxide (Al203), processes 7439-95-4. Magnesium, processes 7446-07-3. Tellurium oxide (TeO2) 7631-86-9. 7789-75-5, Calcium fluoride (CaF2), processes Silica, processes 12064-62-9. Gadolinium oxide (Gd203) 13463-67-7, Titanium oxide (TiO2), processes 65589-84-6. Lithium oxide (Li203) 113482-02-3. Yttrium zirconium oxide (Y0.06Zr0.9702.03) RL: PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process) (in glass-ceramics: aluminate glass ceramic hardness abrasive steel substrate) 12597-68-1, Stainless Steel, uses ΙT

RL: NUU (Other use, unclassified); USES (Uses)

(substrates: aluminate glass ceramic hardness abrasive steel substrate)

L8 MANSWER 2 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2001:726056 CAPLUS

DOCUMENT NUMBER:

136:9674

TITLE:

Equation for estimating the thermal diffusivity. specific heat and thermal conductivity of oxide

glasses

AUTHOR(S):

Inaba, Seiji; Oda, Shuhei; Morinaga, Kenji

CORPORATE SOURCE:

Department of Applied Science for Electronics and Materials. Interdisciplinary Graduate School of Engineering Sciences. Kyushu University. Klasuga.

816-8580, Japan

SOURCE:

Nippon Kinzoku Gakkaishi (2001), 65(8), 680-687

CODEN: NIKGAV: ISSN: 0021-4876

PUBLISHER:

Nippon Kinzoku Gakkai

DOCUMENT TYPE:

Journal

LANGUAGE:

Japanese

AB The thermal diffusivity and sp. heat have been measured at room temperature by laser flash method and AC calorimetry, resp. Thermal conductivity has been determined by the product of d., thermal diffusivity, and sp. heat. The Debye temperature data have been collected from the measurements of ultrasonic method. For the direct calcn. of these properties from glass composition, empirical equations that consist of compositional parameters V1 and G1 were obtained, where V1 is the packing d. parameter and G1 is the dissociation energy of a single component oxide, resp. Exptl. results indicate that the calculated values using our proposed equations are in a good agreement with the measured values in many glasses studied in this work.

IT Borate glasses

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(aluminoborate: empirical equation for estimation of thermal diffusivity.

sp. heat, and thermal conductivity of aluminosilicate, borate, phosphate,

tellurite, and germanate oxide glasses)

IT Phosphate glasses

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(aluminophosphate; empirical equation for estimation of thermal diffusivity.

sp. heat, and thermal conductivity of aluminosilicate, borate, phosphate.

tellurite, and germanate oxide glasses)

IT Heat capacity

Thermal conductivity

Thermal properties

(empirical equation for estimation of thermal diffusivity, sp. heat, and thermal conductivity of aluminosilicate, borate, phosphate, tellurite, and germanate oxide glasses)

IT Aluminosilicate glasses

Borate glasses Glass, properties Phosphate glasses

germanate oxide glasses) 1304-28-5. Baria. properties

ΙT

```
Silicate glasses
    Tellurite glasses
    RL: PRP (Properties); TEM (Technical or engineered material use); USES
        (empirical equation for estimation of thermal diffusivity, sp. heat, and
       thermal conductivity of aluminosilicate, borate, phosphate, tellurite, and
       germanate oxide glasses)
IT
    Glass, properties
    RL: PRP (Properties); TEM (Technical or engineered material use); USES
     (Uses)
        (germanate; empirical equation for estimation of thermal diffusivity, sp.
       heat. and thermal conductivity of aluminosilicate. borate, phosphate,
       tellurite, and germanate oxide glasses)
                                              1308-87-8, Dysprosium
    1303-86-2. Boron sesquioxide, properties
ΙT
    sesquioxide
                  1308-96-9. Europium sesquioxide 1312-81-8, Lanthana
                                      1314-36-9, Yttria, properties
     1313-97-9. Neodymium sesquioxide
    1314-37-0. Ytterbium sesquioxide 12036-32-7, Praseodymium sesquioxide
     12036-40-7. Tantalum sesquioxide 12036-44-1. Thulium sesquioxide
     12055-62-8. Holmium sesquioxide 12060-58-1, Samarium sesquioxide
     12061-16-4, Erbium sesquioxide 12064-62-9, Gadolinium sesquioxide
    RL: PRP (Properties); TEM (Technical or engineered material use); USES
     (Uses)
        (aluminosilicate glass; empirical equation for estimation of thermal
        diffusivity, sp. heat, and thermal conductivity of aluminosilicate, borate.
       phosphate, tellurite, and germanate oxide glasses)
IT
    1304-76-3. Bismuth sesquioxide, properties 1317-36-8, Lead monoxide.
    properties
    RL: PRP (Properties); TEM (Technical or engineered material use); USES
     (Uses)
        (borate glass; empirical equation for estimation of thermal diffusivity, sp.
       heat, and thermal conductivity of aluminosilicate, borate, phosphate,
        tellurite, and germanate oxide glasses)
     1303-28-2. Arsenic pentoxide
                                  1304-56-9. Beryllium monoxide, properties
     1306-19-0. Cadmium monoxide. properties 1306-38-3. Ceria, properties
     1307-96-6. Cobalt monoxide, properties 1308-38-9, Chromia, properties
     1310-53-8. Germanium dioxide. properties
                                               1312-43-2. Indium sesquioxide
     1313-13-9, Manganese dioxide, properties
                                               1314-20-1, Thoria,
                 1314-23-4. Zirconium dioxide, properties
                                                            1314-34-7.
     properties
     Vanadium sesquioxide
                          1314-56-3, Phosphorus pentoxide, properties
     1317-38-0. Copper monoxide, properties 1317-39-1. Copper oxide cu2o.
    properties 7446-07-3, Tellurium dioxide 12024-21-4, Gallium
     sesquioxide
                 12060-08-1, Scandium sesquioxide
                                                    18088-11-4, Rubidium
                 18282-10-5. Tin dioxide
                                           20281-00-9, Cesium oxide cs2o
     oxide rb2o
     20667-12-3. Silver oxide ag2o 21651-19-4. Tin monoxide
    RL: PRP (Properties); TEM (Technical or engineered material use); USES
     (Uses)
        (empirical equation for estimation of thermal diffusivity, sp. heat, and
        thermal conductivity of aluminosilicate, borate, phosphate, tellurite, and
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1305-78-8, Calcia, properties

1309-48-4.

Magnesium monoxide, properties 1313-59-3. Sodium oxide Na2O, properties 1314-11-0, Strontium oxide sro, properties 1344-28-1, Alumina, properties 7631-86-9, Silica, properties 12136-45-7, Potassium oxide k2o, properties 65589-84-6, Lithium oxide Li2O3 RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(oxide glass; empirical equation for estimation of thermal diffusivity, sp. heat, and thermal conductivity of aluminosilicate, borate, phosphate, tellurite, and germanate oxide glasses)

IT 1309-37-1. Iron sesquioxide, properties 1309-64-4, Antimony sesquioxide, properties 1313-27-5, Molybdenum trioxide, properties 1314-13-2, Zinc monoxide, properties 1314-35-8, Tungsten trioxide, properties 12059-63-1, Niobium sesquioxide 13463-67-7, Titania, properties RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(tellurite glass; empirical equation for estimation of thermal diffusivity, sp. heat, and thermal conductivity of aluminosilicate, borate, phosphate, tellurite, and germanate oxide glasses)

**E8 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2001:636306 CAPLUS

DOCUMENT NUMBER: 135:203318

TITLE: Method for single crystal growth of perovskite oxides

INVENTOR(S): Lee, Ho-Yong; Kim, Jae-Suk; Lee, Jong-Bong; Hur,

Tae-Moo; Kim, Doe-Yeon; Hwang, Nong-Moon

PATENT ASSIGNEE(S): Ceracomp Co., Ltd., S. Korea

SOURCE: PCT Int. Appl., 50 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

| PATENT NO. | KIND DATE | APPLICATION NO. | DATE |
|--------------------------------|----------------------------|--|--------------------------|
| WO 2001063021 | A1 20010830 | WO 2001-KR267 | 20010222 |
| | | BA, BB, BG, BR, BY, BEE, ES, FI, GB, GD, G | |
| | | KG, KR, KZ, LC, LK, LI MX, MZ, NO, NZ, PL, P | |
| SE, SG, SI | | | |
| | | SL, SZ, TZ, UG, ZW, A' IE, IT, LU, MC, NL, P' | |
| | | GW. ML. MR. NE. SN. TI | |
| AU 2001036169 JP 2003523919 | A5 20010903 T2 20030812 | AU 2001-36169 JP 2001-561824 | 20010222 20010222 |
| US 2002179000 | A1 20021205 | | 20010611 |
| PRIORITY APPLN. INFO.: | | KR 2000-8916 KR 2001-8685 | A 20000223 A 20010221 |
| | | WO 2001-KR267 | W 20010222 |

AB The invention relates to a method for growing single crystals of perovskite oxides. The method is characterized by comprising the steps of (a) contacting a perovskite seed single with a Perovskite polycrystal and (b) heating the contacted crystals to grow the same structure as the single crystal into the polycrystal, the heating is controlled under conditions which abnormal grains growth is induced in the contacted portion while repressed in the inside of the polycrystal. The method for growing single crystals of perovskite oxides according to this invention has an advantage to provide an effective low cost in manufacturing process for single crystals by using usual heat-treatment process without special equipments. The method for growing single crystals of perovskite oxides according to this invention can be also applicable to other material systems showing abnormal grain growth behavior.

IT Grain growth

(abnormal; method for single crystal growth of perovskite oxides including step of heating combination of seed single crystal and polycrystal under condition of promoting abnormal grain growth)

IT Perovskite-type crystals

(method for single crystal growth of perovskite oxides)

IT Crystal growth

ΙT

(method for single crystal growth of perovskite oxides by changing component ratio)

IT 12047-27-7, Barium titanium oxide (BaTiO3), processes 12060-00-3D, Lead titanium oxide (PbTiO3), solid solution with lead magnesium niobium oxide 12626-81-2, PZT

RL: PEP (Physical, engineering or chemical process); PROC (Process) (method for single crystal growth of perovskite oxides)

1304-28-5, Barium oxide (BaO), processes 1304-76-3, Bismuth oxide (Bi203), processes 1305-78-8, Calcium oxide (CaO), processes 1306-19-0. Cadmium oxide (CdO), processes 1306-38-3, Cerium oxide 1307-96-6. Cobaltous oxide, processes 1308-38-9, (CeO2), processes 1309-37-1, Iron oxide (Fe203). Chromium oxide (Cr203), processes 1309-48-4, Magnesium oxide (MgO), processes 1313-13-9, Manganese oxide, processes Lanthanum oxide (La203) 1313-59-3, Sodium oxide, processes 1313-96-8, Niobium oxide (Nb205) 1313-97-9. Neodymium oxide (nd2o3) 1313-99-1, Nickel oxide (NiO), processes 1314-11-0, Strontium oxide (SrO), processes 1314-13-2, Zinc 1314-23-4, Zirconium oxide (ZrO2), processes oxide (ZnO), processes 1314-36-9, Yttrium oxide (Y2O3), processes 1314-61-0. Tantalum oxide 1317-36-8, Lead oxide (PbO), processes 1344-57-6. Uranium 12055-23-1. Hafnium oxide (HfO2) 12060-08-1. oxide (UO2), processes 12136-45-7, Potassium oxide (K20), processes Scandium oxide (Sc203) 12274-26-9. Samarium oxide (SmO2) 13463-67-7, Titanium oxide (TiO2), 18282-10-5. Tin oxide (SnO2) processes

RL: MOA (Modifier or additive use): PEP (Physical, engineering or chemical process); PROC (Process): USES (Uses)

(method for single crystal growth of perovskite oxides by adjoining seed single crystal to perovskite polycrystal having additives of) 12057-57-7D, Lead magnesium niobium oxide (PbMg0.33Nb0.6703), solid solution with lead titanium oxide

RL: PEP (Physical, engineering or chemical process); PROC (Process) (method for single crystal growth of perovskite oxides by changing component ratio)

1303-86-2. Boron oxide (B203), processes 1310-53-8. Germanium oxide (GeO2), processes 1314-56-3. Phosphorus oxide (P2O5), processes 1314-62-1. Vanadium oxide (V205), processes 1317-38-0, Cupric oxide. processes 1344-28-1, Aluminum oxide (Al203), processes 7631-86-9. Silicon oxide (SiO2), processes 65589-84-6, Lithium oxide (Li203)

RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(method for single crystal growth of perovskite oxides including step of heating combination of seed single crystal and polycrystal under condition of promoting abnormal grain growth by adding)

REFERENCE COUNT:

THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS 5 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

C8 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2001:64217 CAPLUS

DOCUMENT NUMBER:

134:123927

TITLE:

Method for single crystal growth of barium titanate

and barium titanate solid solution

INVENTOR(S):

Lee. Ho-Yong: Kim. Jae-Suk; Lee, Jong-Bong; Hur, Tae-Moo: Kim. Doe-Yeon: Hwang, Nong-Moon: Lee. Byoung-Ki; Chung, Sung Yoon; Kang, Suk-Joong L.

Ceracomp Co., Ltd., S. Korea

PATENT ASSIGNEE(S):

PCT Int. Appl., 58 pp.

SOURCE:

DOCUMENT TYPE:

CODEN: PIXXD2

Patent English

LANGUAGE:

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

| PATENT NO. | KIND DATE | APPLICATION NO. | DATE |
|-----------------|-----------------|---------------------|-----------------|
| WO 2001006042 | A1 20010125 | WO 2000-KR656 | 20000621 |
| | | BA, BB, BG, BR, BY, | |
| CR, CU, CZ, | DE, DK, DM, DZ, | EE, ES, FI, GB, GD, | GE, GH, GM, HR, |
| HU, ID, IL. | IN. IS. JP. KE. | KG, KP, KZ, LC, LK. | LR. LS. LT. LU. |
| LV. MA, MD, | MG. MK. MN. MW. | MX, MZ, NO, NZ, PL. | PT. RO, RU, SD. |
| SE, SG, SI | | | |
| RW: GH, GM, KE, | LS. MW. MZ. SD, | SL. SZ. TZ, UG. ZW, | AT, BE, CH, CY. |
| | | IE, IT, LU, MC, NL, | |
| | | ML, MR, NE, SN, TD, | TG |
| | | KR 1999-23660 | 19990623 |
| AU 2000055747 | A5 20010205 | AU 2000-55747 | 20000621 |
| | | JP 2001-511247 | 20000621 |
| JP 3507821 | B2 20040315 | | |
| US 6482259 | B1 20021119 | US 2001-646610 | |
| US 2003015130 | A1 20030123 | US 2002-163526 | 20020607 |

B2 20040706 US 6758898 A 19990623 PRIORITY APPLN. INFO.: KR 1999-23660 A 20000223 KR 2000-8916 KR 2000-8917 Α 20000223 KR 2000-20620 A 20000419 WO 2000-KR656 W 20000621 US 2001-646610 A1 20010220

AB A method is given for growing single crystals of BaTiO3 and Ba titanate solid solns. [(BaxM1-x)(TiyN1-y)O3]. A method for growing single crystals of BaTiO3 and Ba titanate solid solns. showing the primary and secondary abnormal grain growths with increasing temperature above the liquid formation temperature is characterized by comprising the step for a few secondary abnormal grains to continue to grow at a temperature slightly below the critical temperature where

the secondary abnormal grain growth starts to occur. The method has the advantage to provide an effective low cost in manufacturing process for single crystals by using usual heat-treatment process without special equipment. The method can be also applicable to other material systems showing abnormal grain growth behavior.

IT Heat treatment

(crystal growth of barium titanate and barium titanate solid solns. showing abnormal grain growth behavior by)

IT Grain growth

(crystal growth of barium titanate and barium titanate solid solns. showing abnormal grain growth behavior by heat treatment)

IT Crystal growth

(of barium titanate and barium titanate solid solns. showing abnormal grain growth behavior by heat treatment)

1303-86-2, Boron oxide, processes 1304-28-5, Barium oxide, processes ΙT 1304-76-3. Bismuth oxide bi2o3, processes 1305-78-8. Calcium oxide, 1306-19-0, Cadmium oxide, processes 1306-38-3, Cerium dioxide, processes 1307-96-6. Cobalt oxide coo, processes 1308-38-9. 1309-37-1. Iron oxide fe2o3, processes Chromium oxide cr2o3, processes 1310-53-8. Germanium dioxide. 1309-48-4. Magnesium oxide, processes 1312-81-8, Lanthanum sesquioxide 1313-13-9. processes Manganese dioxide, processes 1313-59-3. Sodium oxide, processes 1313-96-8, Niobium pentoxide 1313-97-9. Neodymium sesquioxide 1313-99-1, Nickel oxide nio, processes 1314-11-0. Strontium oxide, 1314-23-4, Zirconium 1314-13-2. Zinc oxide, processes dioxide, processes 1314-36-9. Yttrium sesquioxide, processes 1314-56-3. Phosphorus pentoxide, processes 1314-61-0. Tantalum pentoxide 1314-62-1, Vanadium pentoxide, processes 1317-36-8. Lead oxide pbo. 1317-38-0. Copper oxide cuo, processes 1344-28-1, Alumina. processes processes 1344-43-0. Manganese oxide mno, processes 7631-86-9. Silica, processes 1344-57-6. Uranium dioxide, processes 12055-23-1, Hafnium dioxide 12060-08-1, Scandium sesquioxide 12136-45-7. Potassium oxide, processes 12274-26-9. Samarium dioxide 13463-67-7. Titanium dioxide, processes 18282-10-5. Tin dioxide **65589-84-6**, Lithium oxide 1i2o3 148070-59-1, Barium titanium

oxide silicate (Ba0.8Ti0.802(Si04)0.2)

RL: MOA (Modifier or additive use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(crystal growth of barium titanate and barium titanate solid solns. showing abnormal grain growth behavior by heat treatment using additive of)

IT 12049-50-2, Calcium titanate 12060-59-2, Strontium titanate RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(crystal growth of barium titanate and barium titanate solid solns. showing abnormal grain growth behavior by heat treatment using seed crystal of)

IT 12047-27-7. Barium titanium oxide batio3, processes 12047-27-7D. Barium titanium oxide batio3, solid solns.

RL: PEP (Physical, engineering or chemical process); PROC (Process) (crystal growth showing abnormal grain growth behavior by heat treatment)

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

(L8_ANSWER_5_OF_7_/ CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1997:666855 CAPLUS

DOCUMENT NUMBER:

127:354580

TITLE:

Cu-Mg-Zn-based magnetic materials with good

temperature characteristics

INVENTOR(S):
PATENT ASSIGNEE(S):

Kang, Sang Won; Park, Jong Hak Samsung Corning Co.,Ltd., S. Korea

SOURCE:

Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------------------|---------|--------------|--------------------|----------------|
| | | | | |
| JP 09263442 | A2 | 19971007 | JP 1996-97661 | 19960327 |
| PRIORITY APPLN. INFO.: | | | JP 1996-97661 | 19960327 |
| AR The materials comp | rise fe | rrites consi | sting of CuO 2 5-9 | MaO 18-24 Fe26 |

AB The materials comprise ferrites consisting of CuO 2.5-9. MgO 18-24, Fe2O3 48-50 mol%, and balance ZnO: whose Fe2O3 component is substituted with 0.1-0.9 mol% Cr2O3 and to which 0.01-0.1 weight% Bi2O3 and 0.1-0.6 weight% Li2O3 are added. The materials also show high initial permeability and low magnetic loss, and are especially useful for rotary transformer cores.

IT Ferrites

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(Cu-Mg-Zn-based ferrite magnetic materials with good temperature characteristics)

IT Magnetic cores

(Cu-Mg-Zn-based ferrite magnetic materials with good temperature characteristics for)

Transformers

(cores: Cu-Mq-Zn-based ferrite magnetic materials with good temperature characteristics for)

198141-40-1P

RL: PNU (Preparation, unclassified); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(Cu-Mg-Zn-based ferrite magnetic materials with good temperature characteristics)

1309-37-1, Iron oxide (Fe203), properties 1309-48-4, Magnesia, IT properties 1314-13-2. Zinc oxide, properties 1317-38-0. Cupric oxide. properties

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(Cu-Mg-Zn-based ferrite magnetic materials with good temperature characteristics)

1304-76-3. Bismuth oxide, properties 1305-78-8, Calcia, properties 1344-43-0, **Manganese** oxide (MnO), properties 7631-86-9.

Silica, properties 65589-84-6, Lithium oxide (Li203)

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(ferrites containing; Cu-Mg-Zn-based ferrite magnetic materials with good temperature characteristics)

1308-38-9. Chromium oxide. properties

RL: PRP (Properties): TEM (Technical or engineered material use): USES (Uses)

(ferrites substituted with: Cu-Mg-Zn-based ferrite magnetic materials with good temperature characteristics)

(L8_ANSWER_6_OF_7_CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1991:438707 CAPLUS

DOCUMENT NUMBER:

115:38707

TITLE:

Electrochromic device containing nickel oxide counter

electrode

Nagai, Junichi: Seike, Tetsuya INVENTOR(S): Asahi Glass Co., Ltd., Japan PATENT ASSIGNEE(S): Jpn. Kokai Tokkyo Koho, 4 pp. SOURCE:

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------------------|------|----------|-----------------|----------|
| | | | | |
| JP 02226122 | A2 | 19900907 | JP 1989-45101 | 19890228 |
| JP 2730139 | B2 | 19980325 | | |
| PRIORITY APPLN. INFO.: | | | JP 1989-45101 | 19890228 |

In the title device comprising an electrode and a counter electrode formed on substrates, an electrochem, coloring substance, and an electrolyte, the counter electrode contains NiOx(x = 0.5-2) and the electrolyte is made by

dissolving an alkali metal salt M+X- (M = Li, Na, K; X = ClO4, BF4, CF3SO3, AsF6, PF6) in propylene carbonate, δ -butyrolactone, sulfolane, or 3-methyl-sulfolane.

IT Optical imaging devices

(electrochromic, containing nickel oxide counter electrodes)

IT 111706-41-3, Nickel oxide (NiO1.5) 134831-06-4, Nickel oxide (NiO0.8) RL: USES (Uses)

(electrochromic devices containing counter electrodes of)

IT 96-48-0 108-32-7. Propylene carbonate 126-33-0, Sulfolane 872-93-5. 3-Methylsulfolane 2926-30-9 3811-04-9. Potassium chlorate 13453-71-9, Lithium chlorate 14283-07-9. Lithium tetrafluoroborate 17029-22-0, Potassium hexafluoroarsenate 21324-40-3, Lithium hexafluorophosphate

RL: USES (Uses)

(electrochromic devices with electrolytes containing)

IT 1306-38-3. Cerium dioxide, uses and miscellaneous 1307-96-6. Cobalt monoxide, uses and miscellaneous 1312-43-2. Indium trioxide 1312-81-8. Lanthanum trioxide 1313-13-9. Manganese dioxide, uses and miscellaneous 1333-82-0. Chromium trioxide 1345-25-1. Ferrous oxide, uses and miscellaneous 12020-60-9. Europium monoxide 12030-49-8. Iridium dioxide 18282-10-5. Tin dioxide 65589-84-6. Lithium oxide (Li203)

RL: USES (Uses)

(electrochromic devices with nickel oxide counter electrodes doped by)

LANSWER 7- OF 71 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1981:524960 CAPLUS

DOCUMENT NUMBER:

95:124960

TITLE:

Grain boundary insulated semiconductor ceramic Matsushita Electric Industrial Co., Ltd., Japan

PATENT ASSIGNEE(S):

Jpn. Kokai Tokkyo Koho. 5 pp.

SOURCE: Jpn. Kokai To CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------------------|------|----------|-----------------|----------|
| | | | | |
| JP 56083919 | A2 | 19810708 | JP 1979-161778 | 19791212 |
| JP 61050371 | B4 | 19861104 | | |
| PRIORITY APPLN. INFO.: | | | JP 1979-161778 | 19791212 |

AB A grain-boundary-insulated semiconductor ceramic is obtained by thermally diffusing an agent from Bi2O3 and Cu2O into grain boundaries of Sr titanate niobate. The diffusion agent may contain MnO2, B2O3, or Li2O3.

IT Semiconductor materials

(ceramic. from strontium niobate titanate containing oxides, grain-boundary-insulated)

IT Transition metal oxides

RL: USES (Uses)

(semiconductor ceramic from strontium niobate titanate containing)

IT 1303-86-2, uses and miscellaneous 1304-76-3, uses and miscellaneous 1313-13-9, uses and miscellaneous 65589-84-6

RL: USES (Uses)

(semiconductor ceramic from strontium niobate titanate containing)

IT 12060-59-2D, solid solution with strontium niobate 12673-59-5D, solid solution with strontium titanate

RL: USES (Uses)

(semiconductor ceramic from, grain-boundary-insulated)

554-13-2. Lithium carbonate 1309-48-4. Magnesium oxide. processes 1313-13-9. Manganese dioxide. processes 7790-69-4. Lithium nitrate 10377-60-3. Magnesium nitrate 10377-66-9. Manganese nitrate RL: PEP (Physical, engineering or chemical process); PROC (Process) (manufacture of secondary Li battery cathodes containing Li Mn Mg oxide)

L5 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2000:754409 CAPLUS

DOCUMENT NUMBER:

133:298835

TITLE:

Preparation of lithium manganese oxides by a

heat treatment for battery use

INVENTOR(S):

Horne, Craig R.; Kumar, Sujeet; Reitz, Hariklia Dris;

Gardner, James T.; Bi, Xiangxin

PATENT ASSIGNEE(S):

NanoGram Corporation, USA

SOURCE:

U.S., 42 pp., Cont.-in-part of U.S. Ser. No. 188,768.

CODEN: USXXAM

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 23

PATENT INFORMATION:

| PAT | ENT NO. | | KIND | DATE | APPLICATION NO. | DATE |
|----------|--------------------------|---------|--------|-----------|---------------------|-----------------|
| US | 6136287 | | A | 20001024 | US 1998-203414 | 19981202 |
| | 6607706 | | B1 | 20030819 | US 1998-188768 | 19981109 |
| WO | 200002775 | 54 | A1 | 20000518 | WO 1999-US26343 | 19991108 |
| | W: CA, RW: AT, PT, | BE, CH, | | , DK, ES, | FI, FR, GB, GR, IE, | IT, LU, MC, NL, |
| EP | 1165442 | | A1 | 20020102 | EP 1999-957527 | 19991108 |
| | R: AT, IE, | | DE, DK | , ES, FR, | GB, GR, IT, LI, LU, | NL, SE, MC, PT, |
| JP : | 200252935 | 52 | T2 | 20020910 | JP 2000-580940 | 19991108 |
| PRIORITY | APPLN.] | INFO.: | | | US 1998-188768 | A2 19981109 |
| | | | | | US 1998-188770 | A 19981109 |
| | | | | | US 1998-203414 | A 19981202 |
| | | | | | US 1999-334203 | A 19990616 |
| | | | | | WO 1999-US26343 | W 19991108 |
| 40 1 1 1 | | | | | | |

- AB Lithium manganese oxide particles have been produced with an average diameter less than about 250 nm. The particles have a high degree of uniformity. The particles are formed by the heat treatment of nanoparticles of manganese oxide. The lithium manganese oxide particles are useful as active materials in the pos. electrodes of lithium based batteries. Improved batteries result from the use of the uniform nanoscale lithium manganese oxide particles.
- IT Secondary batteries

(lithium; preparation of lithium manganese oxides by heat treatment for battery use)

IT Thermal decomposition

(photo-: preparation of lithium manganese oxides by heat treatment for

battery use)

IT Battery cathodes

Heat treatment

(preparation of lithium manganese oxides by heat treatment for battery use)

IT 67-63-0, Isopropyl alcohol, uses

RL: TEM (Technical or engineered material use); USES (Uses)
(IR absorber; preparation of lithium manganese oxides by heat treatment for battery use)

IT 74-85-1. Ethylene, uses

RL: TEM (Technical or engineered material use): USES (Uses)
(lase absorbing gas: preparation of lithium manganese oxides by heat treatment for battery use)

IT 1344-43-0, Manganese oxide mno, reactions 7447-41-8, Lithium chloride, reactions 7773-01-5, Manganese chloride mncl2 7790-69-4, Lithium nitrate 10170-69-1, Manganese carbonyl Mn2(CO)10 10377-66-9, Manganese nitrate

RL: RCT (Reactant); RACT (Reactant or reagent)
(precursor; preparation of lithium manganese oxides by heat treatment for battery use)

IT 96-49-1, Ethylene carbonate 616-38-6, Dimethyl carbonate 1317-34-6, Manganese oxide mn2o3 1317-35-7, Manganese oxide mn3o4 21324-40-3. Lithium hexafluorophosphate

RL: DEV (Device component use); USES (Uses)

(preparation of lithium manganese oxides by heat treatment for battery use)

IT 7439-93-2. Lithium, uses

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(preparation of lithium manganese oxides by heat treatment for battery use)

IT 12057-17-9P. Lithium manganese oxide limn2o4 39457-42-6P.

Lithium manganese oxide 127575-11-5P. Lithium manganese oxide Li2Mn409 301319-26-6P. Lithium manganese oxide (LiMn203.8)

RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(preparation of lithium manganese oxides by heat treatment for battery use) 1313-12-8P. Manganosite

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of lithium manganese oxides by heat treatment for battery use) 7782-44-7. Oxygen. uses

RL: RCT (Reactant); TEM (Technical or engineered material use); RACT (Reactant or reagent); USES (Uses)

(preparation of lithium manganese oxides by heat treatment for battery use)

IT 7440-37-1, Argon, uses 7647-01-0, Hydrochloric acid, uses

·RL: TEM (Technical or engineered material use); USES (Uses)

(preparation of lithium manganese oxides by heat treatment for battery use)
REFERENCE COUNT: 35 THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN ACCESSION NUMBER: 1996:274822 CAPLUS

DOCUMENT NUMBER:

124:321354

TITLE:

Preparation of cathode materials for lithium batteries by melt impregnation method. IV. Preparation of stoichiometric spinel LiMn204 and nonstoichiometric spinel LixMn2O4 and their

electrochemical behavior

AUTHOR(S):

Yoshio, Masaki: Noguchi, Hideyuki; Xia, Yongyao;

Ikeda, Kazutaka

CORPORATE SOURCE:

Dep. Appl. Chem., Fac. Sci. Eng., Saga Univ., Saga,

840. Japan

SOURCE:

Denki Kagaku oyobi Kogyo Butsuri Kagaku (1996), 64(5),

388-393

CODEN: DKOKAZ: ISSN: 0366-9297

PUBLISHER:

Denki Kagaku Kyokai

DOCUMENT TYPE:

Journal

LANGUAGE:

Japanese

The spinel LixMn2Oz (z>4) has been prepared from MnO2 and LiNO3 (or LiOH) under N2 or air by melt impregnation method. Nitrogen gas accelerates the release of excess oxygen from oxygen excess LiMn2Oz, causing the formation of LiMn2O4. Stoichiometric spinel LiMn204 with initial capacity of 130 mAh \cdot g-1 at c.d. 0.4 mA·cm-2 is prepared from IC 12 (International Common MnO2 No.12)-LiOH by two step heating, 470°, 12 h in air and 700°. 24 h in N2. It would be gradually transformed during the cycling to new compds. with the capacity of ca. 110-120 mAh·g-1, which is higher than that of oxygen rich spinel Li1.04Mn2O4.14 with excellent cyclability. The Li1.04Mn204.14 shows one quasi-S type discharge curve. Its electrochem. reaction consisting of two homogeneous reactions was nearly same potentials with those of LiMn204 with a homogeneous and a heterogeneous reactions. Excellent cyclability of it would be due to the continuous and small cell volume change during the cycling.

ΙT Cathodes

> (battery, preparation of stoichiometric spinel LiMn204 and nonstoichiometric spinel LixMn2O4 and their electrochem. behavior)

12057-17-9P, Lithium manganese oxide LiMn204 39457-42-6P. Lithium manganese oxide 171088-91-8DP, Lithium manganese oxide Lil.04Mn204, oxygen-excess 176500-67-7DP, Lithium manganese oxide (Li0.24Mn204), oxygen-excess 176500-68-8DP, Lithium manganese oxide (Li0.34Mn2O4), oxygen-excess 176500-69-9DP. Lithium manganese oxide (Li0.44Mn2O4), oxygen-excess 176500-70-2DP. Lithium manganese oxide (Li0.49Mn204), oxygen 176500-71-3DP. Lithium manganese oxide (Li0.59Mn204). oxygen-excess 177323-75-0DP, Lithium manganese oxide (Li0.79Mn204), oxygen-excess RL: DEV (Device component use); PRP (Properties); SPN (Synthetic

preparation); PREP (Preparation); USES (Uses)

(preparation of stoichiometric spinel LiMn204 and nonstoichiometric spinel

LixMn204 and their electrochem. behavior)

ACCESSION NUMBER:

1991:646496 CAPLUS

DOCUMENT NUMBER:

115:246496

TITLE:

The use of acetates as precursors for the

low-temperature synthesis of lithium manganese oxide

(LiMn204) and lithium cobalt oxide (LiCoO2)

intercalation compounds

AUTHOR(S):

Barboux, P.; Tarascon, J. M.; Shokoohi, F. K.

CORPORATE SOURCE:

Bellcore, Red Bank, NJ, 07701, USA

SOURCE:

Journal of Solid State Chemistry (1991), 94(1), 185-96

CODEN: JSSCBI; ISSN: 0022-4596

DOCUMENT TYPE:

Journal

LANGUAGE:

English

AB The spinel LiMn204 and layered LiM02 (M = Co, Ni) compds., which are of potential interest for Li intercalation applications, have been synthesized at low temps. from aqueous solns. of hydroxides, nitrate, and acetate salts. These phases can be prepared in bulk or thick-film form with their crystallization temperature strongly dependent on O pressure, annealing time, and Li/transition metal ratio. For instance, we succeeded in preparing the spinel LiMn204 at temps. as low as 300°, whereas a temperature of 500° was needed to obtain LiCoO2. Rechargeable Li batteries using the LiMn204 powders, synthesized at low temps., show good capacity and good intercalation cycling behavior. Larger amts. of Li cannot be directly introduced into the spinel phase by heat treatment under less oxidizing atmospheres obtained by use of mixts. of argon and oxygen.

IT Cathodes

(battery, lithium manganese oxide)

IT Inclusion reaction

(intercalation, of lithium into lithium manganese oxide or sodium into manganese sodium oxide)

IT 1344-43-0P. Manganese monoxide, preparation

RL: FORM (Formation, nonpreparative); PREP (Preparation)

(formation of, by thermal decomposition of lithium manganese oxide)

IT 1317-34-6P, Manganese oxide (Mn2O3)

RL: SPN (Synthetic preparation); FORM (Formation, nonpreparative); PREP (Preparation)

(formation of, in sol-gel preparation of lithium manganese oxide spinel)

IT 7439-93-2. Lithium, reactions

RL: RCT (Reactant): RACT (Reactant or reagent)

(intercalation of, into lithium magnesium oxide)

IT 7440-23-5. Sodium, reactions

RL: RCT (Reactant); RACT (Reactant or reagent)

(intercalation of, into manganese sodium oxide)

IT 12057-17-9P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation and thermal decomposition and reversible intercalation of lithium into)

IT 71-48-7. Cobalt(2+) acetate 373-02-4. Nickel(2+) acetate 638-38-0.

Manganese(2+) acetate 10377-66-9, Manganous nitrate

RL: RCT (Reactant); RACT (Reactant or reagent)

(reaction of, with lithium and ammonium hydroxides)

IT 1310-65-2, Lithium hydroxide

RL: RCT (Reactant); RACT (Reactant or reagent)

(reaction of, with transition metal salts and ammonium hydroxide)

IT 137184-05-5, Manganese sodium oxide (MnNa0.2502)

RL: RCT (Reactant); RACT (Reactant or reagent)

(sol-gel preparation and intercalation of sodium into)

IT 12190-79-3P, Cobalt lithium oxide (CoLiO2)

RL: SPN (Synthetic preparation); PREP (Preparation)

(sol-gel preparation of)

IT 137184-04-4, Manganese sodium oxide (MnNa0.4402)

RL: PROC (Process)

(thermal formation of)

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ANSWER 1 OF 1 REGISTRY COPYRIGHT 2004 ACS on STN
L1
    12057-17-9 REGISTRY
RN
CN Lithium manganese oxide (LiMn204) (6CI, 7CI, 9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
   Manganate (Mn2O41-), lithium
OTHER NAMES:
    Lithium manganate (LiMn204)
    Lithium manganese oxide (Li0.5MnO2)
CN
    Lithium manganite (LiMn204)
CN
CN
    LM 4
CN
    LM 4 (metal oxide)
DR
    89412-12-4
MF
    Li . Mn . 0
ΑF
    Li Mn2 04
CI
    TIS
LC
                 CA. CAOLD, CAPLUS, CHEMCATS, CHEMLIST, CSCHEM, IFICDB.
    STN Files:
       IFIUDB, TOXCENTER, USPAT2, USPATFULL
    Other Sources: TSCA**
         (**Enter CHEMLIST File for up-to-date regulatory information)
DT.CA CAplus document type: Conference: Dissertation; Journal; Patent;
      Preprint
      Roles from patents: ANST (Analytical study); PREP (Preparation); PROC
RL.P
      (Process): PRP (Properties): RACT (Reactant or reagent): USES (Uses)
      Roles for non-specific derivatives from patents: PREP (Preparation);
      PROC (Process); PRP (Properties); USES (Uses)
      Roles from non-patents: ANST (Analytical study); BIOL (Biological
      study); CMBI (Combinatorial study); FORM (Formation, nonpreparative);
      MSC (Miscellaneous); OCCU (Occurrence); PREP (Preparation); PROC
       (Process): PRP (Properties); RACT (Reactant or reagent): USES (Uses):
      NORL (No role in record)
RLD.NP Roles for non-specific derivatives from non-patents: ANST (Analytical
      study): FORM (Formation, nonpreparative); PREP (Preparation); PROC
```

| Component | Ratio | Component Registry Number |
|--------------|---|--|
| ============ | +====================================== | +===================================== |
| 0 | 4 | 17778-80-2 |
| Mn | j 2 | 7439-96-5 |
| Li | 1 | 7439-93-2 |

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2601 REFERENCES IN FILE CA (1907 TO DATE)
78 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
2606 REFERENCES IN FILE CAPLUS (1907 TO DATE)
6 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

(Process); PRP (Properties); RACT (Reactant or reagent); USES (Uses)

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=> d hist
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(FILE 'HOME' ENTERED AT 16:09:18 ON 08 AUG 2004)

FILE 'REGISTRY' ENTERED AT 16:09:28 ON 08 AUG 2004

E LIMN204/MF

L1 1 S E3

FILE 'CAPLUS' ENTERED AT 16:10:15 ON 08 AUG 2004

L2 170 S L1 AND PREPARATION

L3 0 S L2 AND OXYGEN AND INERT L4 27 S L2 AND (AIR OR OXYGEN)

L5 5 S L4 AND (ARGON OR NITROGEN)

=> d ibib ab it 1-

YOU HAVE REQUESTED DATA FROM 5 ANSWERS - CONTINUE? Y/(N):y

L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2001:830604 CAPLUS

DOCUMENT NUMBER:

135:350822

TITLE:

Procedure for the crystallization of a thin film of a

lithium transition metal oxide for a secondary battery

electrode

INVENTOR(S):

Lee, Jai Young; Kang, Youn Seon; Lee, Ho; Park, Sung

Chul; Kang, Yong Mook

PATENT ASSIGNEE(S):

Korea Advanced Institute of Science & Technology.

Daejeon, S. Korea

SOURCE:

Ger. Offen., 10 pp.

CODEN: GWXXBX

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE | |
|------------------------|------|----------|------------------|---------|---|
| | | | | | - |
| DE 10053733 | A1 | 20011115 | DE 2000-10053733 | 2000103 | 0 |
| DE 10053733 | B4 | 20040805 | | | |
| US 6376027 | B1 | 20020423 | US 2000-688987 | 2000101 | 7 |
| PRIORITY APPLN. INFO.: | | | KR 2000-23286 | 2000050 | 1 |

AB A procedure is presented for the crystallization of a thin film of a Li transition metal oxide for an electrode of a secondary Li thin film battery. The thin film of Li transition metal oxide is deposited by 0 or Ar microwave or high-frequency plasma and has a high degree of crystallinity and excellent electrochem. characteristics. The plasma treatment solves, not only the problem of high deposition temps. approaching 750 C and a procedure lasting from several hours to 10 h by the conventional crystallization procedure for thin films of Li transition metal oxides, but also the problem using a glass or a metal substrate with lower m.p.

IT Secondary batteries

. •

(lithium; procedure for crystallization of a thin film of a lithium transition metal oxide for a secondary battery electrode)

IT Transition metal oxides

RL: SPN (Synthetic preparation); TEM (Technical or engineered material

use): PREP (Preparation): USES (Uses)

(lithium: procedure for crystallization of a thin film of a lithium transition metal oxide for a secondary battery electrode)

IT Vapor deposition process

(plasma: procedure for crystallization of a thin film of a lithium transition metal oxide for a secondary battery electrode)

IT Battery electrodes

Crystallization

Film electrodes

(procedure for crystallization of a thin film of a lithium transition metal oxide for a secondary battery electrode)

IT 7439-90-9, Krypton, processes 7440-01-9, Neon, processes 7440-37-1,

Argon, processes 7440-59-7, Helium, processes 7440-63-3,

Xenon, processes 7782-44-7, Oxygen, processes

RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(procedure for crystallization of a thin film of a lithium transition metal oxide for a secondary battery electrode)

IT 1314-62-1P. Vanadium oxide (V205), preparation 10377-52-3P.

Lithium phosphate (Li3PO4) 12024-01-0P, Gallium lithium oxide (GaLiO2) 12031-65-1P, Lithium nickel oxide (LiNiO2) 12037-42-2P, Vanadium oxide

(V6013) 12039-13-3P. Titanium sulfide (TiS2) 12057-17-9P.

Lithium manganese oxide (LiMn204) 12057-19-1P, Lithium titanium oxide (LiTiO2) 12190-79-3P, Cobalt lithium oxide (CoLiO2) 12615-39-3P,

Aluminum 50, lithium 50 (atomic) 18282-10-5P, Tin oxide (SnO2)

RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(procedure for crystallization of a thin film of a lithium transition metal oxide for a secondary battery electrode)

IT 7440-06-4. Platinum, uses 7440-21-3, Silicon, uses 7440-32-6.

Titanium, uses 7631-86-9, Silica, uses

RL: TEM (Technical or engineered material use); USES (Uses)

(substrate; procedure for crystallization of a thin film of a lithium transition metal oxide for a secondary battery electrode)

REFERENCE COUNT:

THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2004 ACS on STN

8

ACCESSION NUMBER:

2001:745667 CAPLUS

DOCUMENT NUMBER:

135:291379

TITLE:

Cathode active materials, their manufacture, and

lithium ion secondary batteries using them

INVENTOR(S):

Nozaki, Ayumi; Maekawa, Takeyuki; Miyashita, Shoji

PATENT ASSIGNEE(S):

Mitsubishi Electric Corp., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 9 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent Japanese

LANGUAGE:

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 2001283847 A2 20011012 JP 2000-90928 20000329
PRIORITY APPLN. INFO.: JP 2000-90928 20000329

Cathode active materials are manufactured by mixing Li inorg. salts, Mn inorg. salts, and Mg inorg. salts, heating the mixts. at $600\text{-}900^\circ$ under oxidizing atmospheric, and heating them at $600\text{-}900^\circ$ under reducing atmospheric Alternatively, cathode active materials are manufactured by forming solns. containing Li ion, Mn ion, and Mg ion at a ratio of x:1-y:y (x = 0.8-1.15, y = 0.01-0.1) and agents capable of coordinating with Li, Mn Mg, removing solvents from the solns., and heating the resulting precursors at $600\text{-}900^\circ$ under reducing atmospheric or inert atmospheric Cathode active materials containing spinel-type oxides Lix(Mn1-yMgy)20z (x = 0.8-1.15, y = 0.01-0.1, z = 4-4.2) and secondary Li batteries using the materials are also claimed. The batteries show high initial irreversible capacity and good durability at high temperature

IT Air

Mg

(heating atmospheric; manufacture of secondary Li battery cathodes containing Li $\,\mathrm{Mn}$

oxide)

IT Secondary batteries

(lithium; manufacture of secondary Li battery cathodes containing Li Mn Mg oxide)

IT Battery cathodes

Firing (heat treating)

(manufacture of secondary Li battery cathodes containing Li Mn Mg oxide)

IT Carboxylic acids, preparation

RL: PEP (Physical, engineering or chemical process); SPN (Synthetic

preparation); PREP (Preparation); PROC (Process)

(salts, cathode precursors; manufacture of secondary Li battery cathodes containing Li Mn Mg oxide)

IT 1333-74-0. Hydrogen, uses 7440-37-1, **Argon**, uses

RL: NUU (Other use, unclassified); USES (Uses)

(heating atmospheric: manufacture of secondary Li battery cathodes containing Li $\mbox{\rm Mn}$

Mg

IT 12057-17-9. Lithium manganese oxide (LiMn204)

RL: DEV (Device component use); USES (Uses)

(manufacture of secondary Li battery cathodes containing Li Mn Mg oxide)

IT 175786-46-6P. Lithium magnesium manganese oxide

RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(manufacture of secondary Li battery cathodes containing Li Mn Mg oxide)

IT 77-92-9. Citric acid, processes 546-93-0, Magnesium carbonate